

### AMENDMENTS TO THE CLAIMS

Please amend Claim 1 as indicated below.

1. **(Currently amended)** A method for detecting an analyte in a sample comprising:

(a) contacting a sample with a fluorophore-labeled aptamer bound to a solid support;

(b) directly illuminating the aptamer with polarized light whereby the direct illumination of the fluorophore directly excites the fluorophore;

(c) measuring the fluorescence anisotropy of the fluorophore when said fluorophore-labeled aptamer is bound to said analyte; and

(d) identifying the presence or amount of the analyte when ~~[[said]]~~ the measured fluorescence anisotropy ~~measurement~~ is greater than an anisotropy measurement obtained in the absence of bound ~~[[the ]]~~ analyte.

2. **(Previously Presented)** The method of claim 1 wherein the solid support is a bead.

3. **(Previously Presented)** The method of claim 2 wherein the bead is a silica bead.

4. **(Previously Presented)** The method of claim 2 wherein the bead has a diameter between about 1  $\mu\text{m}$  and about 10  $\mu\text{m}$ .

5. **(Previously Presented)** The method of claim 4 wherein the bead has a diameter of about 5  $\mu\text{m}$ .

6. **(Previously Presented)** The method of claim 2 wherein the bead is suspended in solution.

7. **(Previously Presented)** The method of claim 2 wherein the bead is arranged in a two-dimensional array.

8. **(Previously Presented)** The method of claim 1 wherein the aptamer comprises between about 10 and about 100 nucleotides.

9. **(Previously Presented)** The method of claim 1 wherein the aptamer is labeled with a fluorophore selected from the group consisting of fluorescein derivatives, eosin derivatives, coumarin derivatives, and rhodamine derivatives.

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10. **(Previously Presented)** The method of claim 9 wherein the aptamer is labeled with carboxyfluorescein (FAM).
11. **(Previously Presented)** The method of claim 1 wherein the aptamer is part of an array of aptamers.
12. **(Previously Presented)** The method of claim 11 wherein the array comprises two or more addressable locations.
13. **(Previously Presented)** The method of claim 12 wherein each addressable location comprises a single type of aptamer.
14. **(Previously Presented)** The method of claim 12 wherein each addressable location comprises multiple types of aptamers.
15. **(Previously Presented)** The method of claim 14 wherein each type of aptamer is labeled with a fluorophore with unique spectral characteristics.
16. **(Previously Presented)** The method of claim 1 wherein the polarized light is laser light.
17. **(Previously Presented)** The method of claim 1 wherein the analyte is associated with a disease or disorder.
18. **(Previously Presented)** The method of claim 1 wherein the sample is obtained from a patient suspected of suffering from a disease or disorder.
19. **(Previously Presented)** The method of claim 1 wherein the analyte is a protein.
20. **(Previously Presented)** The method of claim 1 wherein the analyte is a metabolite.
21. **(Previously Presented)** The method of claim 1 wherein the sample is from a human patient and the analyte is associated with a disease or disorder.